

UNITED STATES PATENT AND TRADEMARK OFFICE



APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/583,342	05/31/2000	Frederic Bushman	1211.002US1	2389
21186 SCHWFGN	7590 06/12/2002 MAN LUNDBERG. WO	ESSNER & KLUTH, P.A.	EXAMI	NER
P.O. BOX 29		CHAKRABARTI, ARUN K		
			ART UNIT	PAPER NUMBER
			1634 DATE MAILED: 06/12/2002	7

Please find below and/or attached an Office communication concerning this application or proceeding.

· · · · · · · · · · · · · · · · · · ·		Application No.	Applicant(s)
		09/583,342	BUSHMAN ET AL.
	Office Action Summary	Examiner	Art Unit
		Arun Chakrabarti	1634
		cation appears on the cover she	et with the correspondence address
Period for	r Reply		
THE N - Exten after S - If the - If NO - Failur	DRTENED STATUTORY PERIOD FOR ALLING DATE OF THIS COMMUNION SIGNATION OF THIS COMMUNION OF THIS COMMUNION OF THE PROPERTY OF THIS COMMUNION OF THE PROPERTY OF THIS COMMUNION OF THE PROPERTY OF THIS COMMUNICATION OF THE CO	CATION. of 37 CFR 1.136(a). In no event, however, r unication. or days, a reply within the statutory minimum tutory period will apply and will expire SIX (6)	nay a reply be timely filed of thirty (30) days will be considered timely.) MONTHS from the mailing date of this communication. one ARANDONED (35 U.S.C. § 133).
1)⊠	Responsive to communication(s) fil	ed on <u>02 May 2002</u> .	
2a)□	This action is FINAL .	2b)⊠ This action is non-final.	
3)	Since this application is in condition closed in accordance with the praction of Claims	n for allowance except for formatice under <i>Ex parte Quayle</i> , 193	al matters, prosecution as to the merits is 35 C.D. 11, 453 O.G. 213.
•	Claim(s) 1-16 and 20 is/are pending	in the application.	
7)63	4a) Of the above claim(s) is/a	re withdrawn from consideratio	n.
	Claim(s) is/are allowed.		
	Claim(s) 1-16 and 20 is/are rejected	l.	
=	Claim(s) is/are objected to.		
8)□	Claim(s) are subject to restrict	ction and/or election requireme	nt.
	ion Papers		
9)[The specification is objected to by th	e Examiner.	
10)	The drawing(s) filed on is/are	a) accepted or b) dobjected to	to by the Examiner.
	Applicant may not request that any ob	jection to the drawing(s) be held in	a abeyance. See 37 CFR 1.85(a).
11)	The proposed drawing correction file		
	If approved, corrected drawings are re		1.
12)	The oath or declaration is objected t	o by the Examiner.	
Priority	under 35 U.S.C. §§ 119 and 120		0.0.0.440(.) (1) 0.7 (5)
	Acknowledgment is made of a clair	n for foreign priority under 35 U	I.S.C. § 119(a)-(d) or (i).
а) ☐ All b) ☐ Some * c) ☐ None of:		
	1. Certified copies of the priority	y documents have been receive	ed.
	2. Certified copies of the priority	y documents have been receive	ed in Application No
*	3. Copies of the certified copies application from the Intel See the attached detailed Office acti	national Bureau (PC) Rule 17	e been received in this National Stage .2(a)). es not received.
14)	Acknowledgment is made of a claim	for domestic priority under 35	U.S.C. § 119(e) (to a provisional application).
ł	a) The translation of the foreign land Acknowledgment is made of a claim.	anguage provisional application	has been received.
Attachme			
1) No	tice of References Cited (PTO-892) tice of Draftsperson's Patent Drawing Review ormation Disclosure Statement(s) (PTO-1449)	(PTO-948) 5) \(\bigcup \)	nterview Summary (PTO-413) Paper No(s) Notice of Informal Patent Application (PTO-152) Other: Detailed Action .

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DETAILED ACTION

Specification

1. Applicant's election of Group I, corresponding to claims 1-16 and 20, in Paper No. 6 is acknowledged. Because applicant did not distinctly and specifically point out the supposed errors in the restriction requirement, the election has been treated as an election without traverse (MPEP § 818.03(a)).

Claim Rejections - 35 USC § 102

2. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371© of this title before the invention thereof by the applicant for patent.

The changes made to 35 U.S.C. 102(e) by the American Inventors Protection Act of 1999 (AIPA) do not apply to the examination of this application as the application being examined was not (1) filed on or after November 29, 2000, or (2) voluntarily published under 35 U.S.C. 122(b). Therefore, this application is examined under 35 U.S.C. 102(e) prior to the amendment by the AIPA (pre-AIPA 35 U.S.C. 102(e)).

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3. Claims 1-16 and 20 are rejected under 35 U.S.C. 102 (e) as being anticipated by Lynch et al. (U.S. Patent 6,197,527 B1) (March 6, 2000).

Lynch et al teach a high-throughput method of screening compounds capable of modulating topoisomerase activity (Abstract and Column 4, lines 11-34) comprising:

- a) incubating at least a first nucleic acid, a topoisomerase and a potential topoisomerase-modulating compound, wherein the nucleic acid comprises at least one tag (Figures 1-4 and Examples 1-3 and Column 19, lines 31-50), and
- b) assaying for nucleic acid religation inherently (Figures 1-4 and Column 19, line 53 to Column 20, line 8). This inherence is deduced from the fact that when a topoisomerase becomes trapped in a covalent intermediate resulting in cleavage of the nucleic acid between the two labels, the quenching is lost and an increase in signal is observed. Naturally, when there is decreased religation which is equivalent to increased cleavage, an increase in signal will be observed. On the other hand, when there is increased religation, a decrease in signal will be observed.

Lynch et al teach a high-throughput method, wherein the nucleic acid is DNA and RNA (Column 8, lines 57-62).

Lynch et al teach a high-throughput method, wherein the at least one tag is a detection tag or an affinity tag (Column 2, lines 33-44 and Column 13, line 24 to Column 14, line 63 and Column 10, lines 23-36).

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Lynch et al teach a high-throughput method, wherein the method comprises incubating at least a first nucleic acid and a second nucleic acid (Column 16, line 67 to column 17, line 10).

Lynch et al teach a high-throughput method, wherein the second nucleic acid is a religation strand comprising oligonucleotides operatively associated with at least one marker tag (Column 16, line 67 to column 17, line 26).

Lynch et al teach a high-throughput method, wherein the first nucleic acid is operatively associated with an affinity tag and the second nucleic acid is operatively associated with a detection tag (Column 16, line 67 to column 17, line 26).

Lynch et al teach a high-throughput method, wherein the assay detects for topoisomerase inhibitors and activators (Column 1, line 66 to column 2, line 2 and Column 4, lines 17-24).

Lynch et al teach a high-throughput method, wherein the topoisomerase is a Type I or Type II or Type III or Type IV isomerase (Column 16, lines 12-44).

Lynch et al teach a high-throughput method, wherein assaying comprises measuring the level of nucleic acid religation activity in the presence and absence of the topoisomerase modulating compound (Figures 1-4 and Column 19, line 53 to Column 20, line 8).

Lynch et al teach a high-throughput method, wherein the level of religation activity is inversely proportional to the effectiveness of the topoisomerase-inhibitory compound (Column 20, lines 4-8).

Lynch et al teach a high-throughput method, wherein step (a) is performed on a solid support (Figures 1, 3 and Column 15, lines 38-65 and Column 17, lines 23-63).

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Lynch et al teach a high-throughput method, wherein step (a) is performed in a liquid phase (Column 17, line 65 to Column 19, line 50).

Lynch et al teach a high-throughput method, wherein the nucleic acid and topoisomerase are covalently complexed, wherein the topoisomerase retains its religation activity (Column 15, lines 11-15 and Figure 1).

Lynch et al teach a kit for screening compounds that modulate topoisomerase religation activity comprising:

- a) a substrate nucleic acid comprising a first tag,
- b) a religation nucleic acid comprising a second tag,
- c) a topoisomerase, and
- d) a means for measuring nucleic acid religation activity of a test mixture comprising a), b) and c) in the presence or absence of a topoisomerase modulating compound (Column 2, lines 20-47 and Column 19, line 53 to Column 20, line 57).

Conclusion

4. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Arun Chakrabarti, Ph.D., whose telephone number is (703) 306-5818. The examiner can normally be reached on 7:00 AM-4:30 PM from Monday to Friday. If attempts to reach the examiner by telephone are unsuccessful, the examiner's

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supervisor, Gary Jones, can be reached on (703) 308-1152. The fax phone number for this Group is (703) 305-7401. Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the Group analyst Chantae Dessau whose telephone number is (703) 605-1237.

Arun Chakrabarti,

Patent Examiner,

May 21, 2002

Supervisory Patent Examiner Technology Center 1600